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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
09/802,520	03/09/2001	Preeti Lal	PC-0037 US	9214
27904 7	590 07/09/2002			
	INCYTE GENOMICS, INC.		EXAMINER	
3160 PORTER PALO ALTO,		DAVIS, MINH TAM B		Н ТАМ В
			ART UNIT	PAPER NUMBER
			1642	01
			DATE MAILED: 07/09/2002	1

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application No.	Applicant(s)		
Office Action Summary		09/802,520	LAL ET AL.		
		Examiner	Art Unit		
	-	MINH-TAM DAVIS	1642		
	The MAILING DATE of this communication app				
Period fo	• •				
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. - If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. - Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). - Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b). Status					
1)[Responsive to communication(s) filed on 29 A	April 2002 .			
2a)□		is action is non-final.			
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.					
Disposition of Claims					
4) Claim(s) 1-20 is/are pending in the application.					
4a) Of the above claim(s) is/are withdrawn from consideration.					
5) Claim(s) is/are allowed.					
6)□	Claim(s) is/are rejected.				
	Claim(s) is/are objected to.				
	Claim(s) <u>1-20</u> are subject to restriction and/or e	election requirement.	,		
Application Papers					
9) The specification is objected to by the Examiner.					
10) The drawing(s) filed on is/are: a) accepted or b) objected to by the Examiner. Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).					
11) The proposed drawing correction filed on is: a) approved b) disapproved by the Examiner.					
If approved, corrected drawings are required in reply to this Office action.					
12)☐ The oath or declaration is objected to by the Examiner.					
Priority under 35 U.S.C. §§ 119 and 120					
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).					
a) All b) Some * c) None of:					
1. Certified copies of the priority documents have been received.					
	2. Certified copies of the priority documents have been received in Application No				
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 					
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).					
a) ☐ The translation of the foreign language provisional application has been received. 15)☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.					
Attachment(s)					
2) D Notice	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	5) Notice of Informal I	v (PTO-413) Paper No(s) Patent Application (PTO-152)		

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DETAILED ACTION

Effective February 7, 1998, the Group Art Unit location has been changed, and the examiner of the application has been changed. To aid in correlating any papers for this application, all further correspondence regarding this application should be directed to Minh-Tam Davis, Group Art Unit 1642.

The text of those sections of Title 35, U.S. Code not included in this action can be found in a prior Office action.

Applicant election of group I, claims 1-6 and amendment of claim 17 in paper No:8, is acknowledged.

After review and reconsideration, claims 1-20 require further restriction.

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

Group I. Claims 1-6, drawn to a nucleic acid sequence of SEQ ID NO:2, or a nucleic acid sequence encoding SEQ ID NO:1, fragments selected from SEQ ID NO:3-9, a vector comprising a nucleic acid encoding SEQ ID NO:1, a host cell comprising said vector, and a method for producing a protein, classified in class 536, subclass 23.1.

Group II. Claims 1-6, drawn to a nucleic acid sequence of which is a variant of SEQ ID NO:2, comprising SEQ ID NO:10, or a complement thereof, a vector comprising SEQ ID NO:10, a host cell comprising said vector, and a method for producing a protein, classified in class 536, subclass 23.1.

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Group III. Claims 7, 9, drawn to a method for detecting the expression of a nucleic acid, using hybridization, classified in class 435, subclass 6.

Group IV. Claim 8, drawn to a method for detecting the expression of a nucleic acid, using amplification, classified in class 435, subclass.

Group V. Claim 10, drawn to a method for detecting prostate hyperplasia, classified in class 435, subclass 6.

Group VI. Claim 10, drawn to a method for detecting prostate cancer, classified in class 435, subclass 6.

Groups VII. Claims 11-12, drawn to a method for screening compounds that bind specifically to a nucleic acid encoding SEQ ID NO:1, classified in class 435, subclass 6.

Group VIII. Claims 13-14, drawn to a protein of SEQ ID NO:1 or fragments thereof, classified in class 530, subclass 350.

Group IX. Claims 15-16, drawn to a method for screening compounds that specifically binds to SEQ ID NO:1, classified in class 435, subclass 7.1.

Group X. Claim 17, drawn to an antibody specific for SEQ ID NO:1 or an antigenic epitope or a biologically active portion thereof, classified in class 435, subclass 387.1

Group XI. Claim 18, drawn to a method for producing said antibody, using SEQ ID NO:1 or an antigenic epitope or a biologically active portion thereof, classified in class 435, subclass 7.1.

Group XII. Claims 19-20, drawn to a method for detecting prostate hyperplasia, using an antibody, classified in class 435, subclass 7.1.

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Group XIII. Claims 19-20, drawn to a method for detecting prostate cancer, using an antibody, classified in class 435, subclass 7.1.

In addition, upon election of group I, further election of the following species is required"

Any one sequence from SEQ ID Nos: 3-9.

Upon election of group VII further election of any of the following species is required:

Any one of the molecules recited in claim 12.

Upon election of group IX, further election of any of the following species is required:

Any one of the molecules recited in claim16

The inventions are distinct, each from each other because of the following reasons:

Inventions (I, II, VIII, X-XI) and (III-VII, IX, XXII-XIII) are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (M.P.E.P. 806.05 (h). In this instant case, a polypeptide could be used for several purposes, e.g. for biochemical assay, for making antibodies, and for making an affinity column to purify its antibodies; a DNA sequence could be used for the detection of similar DNA or RNA sequences, for making an expression vector, and for producing its encoded protein; and an antibody

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could be used for immunoassay, for purification of its antigen, and for detection of diseases.

The products of groups I, II, VIII, X-XI are patentably distinct, because they are drawn to entirely different biochemicals, having different structures, biological properties and activities that are not interchangeable and cannot be used in place of each other.

The methods of groups III-VII, IX, XXII-XIII are distinct from each other because they differ at least in objectives, method steps, reagents and/or dosages, and/or schedules used, response variables and criteria for success.

The species of the fragments of SEQ ID Nos: 3-9 are distinct because they are structurally distinct.

The species of claim 12, DNA molecules, RNA molecules, peptide nucleic acids, artificial chromosome constructions, peptides, transcription factors, repressors, and regulatory molecules, wherein regulatory molecules are generic to transcription factors and repressors are distinct, because they are structurally distinct, having different characteristics and properties.

The species of claim 16, 1) DNA molecules, 2) RNA molecules, 3) peptide nucleic acids, 4) peptides, 5) proteins, 6) mimetics, 7) agonists, antibodies or immunoglobulins, antagonists, or inhibitors, and 9) drugs are distinct, because they are structurally distinct, having different characteristics and properties.

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Because these inventions are distinct for the reason given above and have acquired a separate status in the art, and because the searches for the groups are not co-extensive, restriction for examination purposes as indicated is proper.

Applicants are required under 35 USC 121 to elect a single disclosed group for prosecution on the merits to which the claims shall be restricted. Applicant is further advised that if Applicant elects a group having species requirement, a response to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 USC 103 of the other invention.

Applicants are reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 C.F.R. 1.48(b) if

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one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendement of inventorship must be accompanied by a diligently-filed petition under 37 C.F.R. 1.48(b) and by the fee required under 37 C.F.R. 1.17(h).

Any inquiry concerning this communication or earlier communications from the examiner should be directed to MINH-TAM DAVIS whose telephone number is 703-305-2008. The examiner can normally be reached on 9:30AM-4:00PM.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, ANTHONY CAPUTA can be reached on 703-308-3995. The fax phone numbers for the organization where this application or proceeding is assigned are 703-872-9306 for regular communications and 703-872-9307 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-0916.

MINH TAM DAVIS

April 05/2002